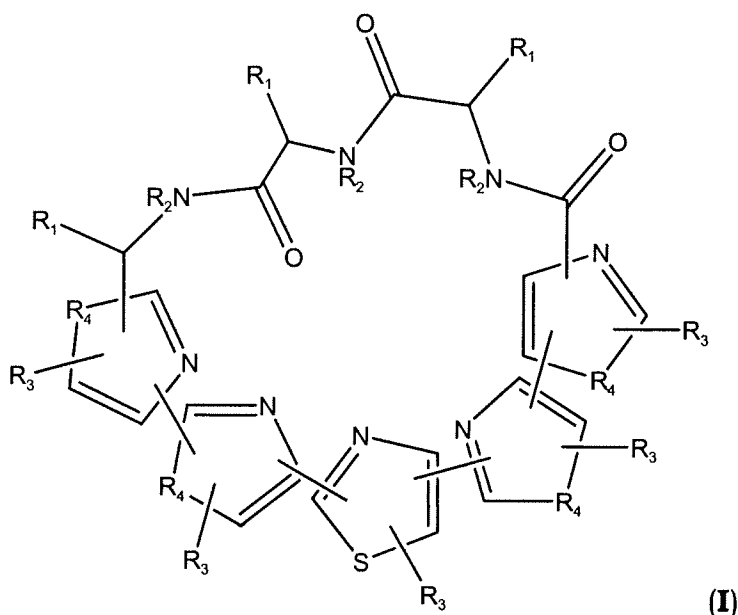


This listing of claims replaces all prior versions and listings of claims in the application:

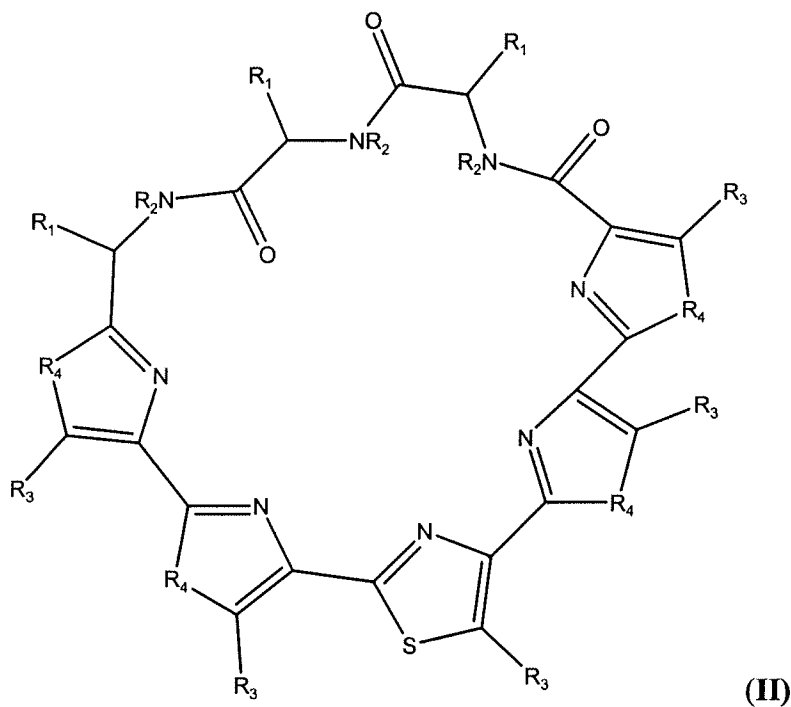
1. (Currently Amended) A purified compound of general formula I:



wherein R<sub>1</sub> are each independently selected from the group consisting of hydrogen, halogen, cyano, hydroxyl, nitro, azido, substituted or unsubstituted alkyl, substituted or unsubstituted alkylidene, substituted or unsubstituted alkenyl, substituted or unsubstituted alkynyl, substituted or unsubstituted alkoxy, substituted or unsubstituted aryl, substituted or unsubstituted heterocyclic group and substituted or unsubstituted acyl; R<sub>3</sub> groups are each independently selected from the group consisting of hydrogen, halogen, cyano, hydroxyl, nitro, azido, substituted or unsubstituted alkyl, substituted or unsubstituted alkenyl, substituted or unsubstituted alkynyl, substituted or unsubstituted alkoxy, substituted or unsubstituted aryl, substituted or unsubstituted heterocyclic group and substituted or unsubstituted acyl

R<sub>4</sub> groups are each independently selected from NR<sub>2</sub>, O and S; and R<sub>2</sub> groups are each independently selected from the group consisting of hydrogen, substituted or unsubstituted alkyl, substituted or unsubstituted aryl, substituted or unsubstituted alkoxy and substituted or unsubstituted acyl, or a pharmaceutically acceptable salt, derivative, prodrug or stereoisomer thereof.

2. (Original) The compound according to claim 1, having the following formula **II**:



wherein R<sub>1</sub>, R<sub>2</sub>, R<sub>3</sub> and R<sub>4</sub> are as defined in claim 1.

3. (Currently Amended) The compound according to ~~claims~~ claim 1 or 2, wherein R<sub>1</sub> are each independently selected from substituted or unsubstituted alkyl and substituted or unsubstituted alkylidene.

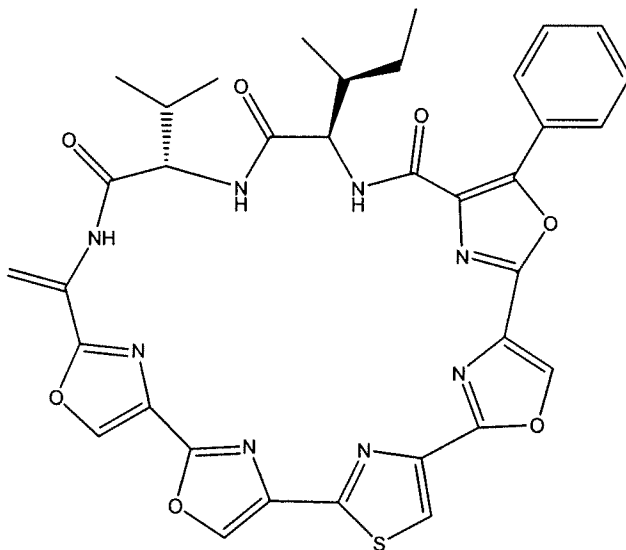
4. (Currently Amended) The compound according to ~~any of claims~~ claim 1 to 3, wherein R<sub>2</sub> are each independently selected from H and substituted or unsubstituted alkyl.

6. (Previously Presented) The compound according to claim 1, wherein R<sub>4</sub> are each O.

CC(C)C(C)C(=O)NC(C)C(=O)NC(C)C(=O)Nc1c(Cc2ccccc2)oc3cc(C4C=CC5=C4N=C5C6C=CC7=C6N=C7C8C=CC9=C8N=C9C)nn31

or a pharmaceutically acceptable salt, derivative, prodrug or stereoisomer thereof.

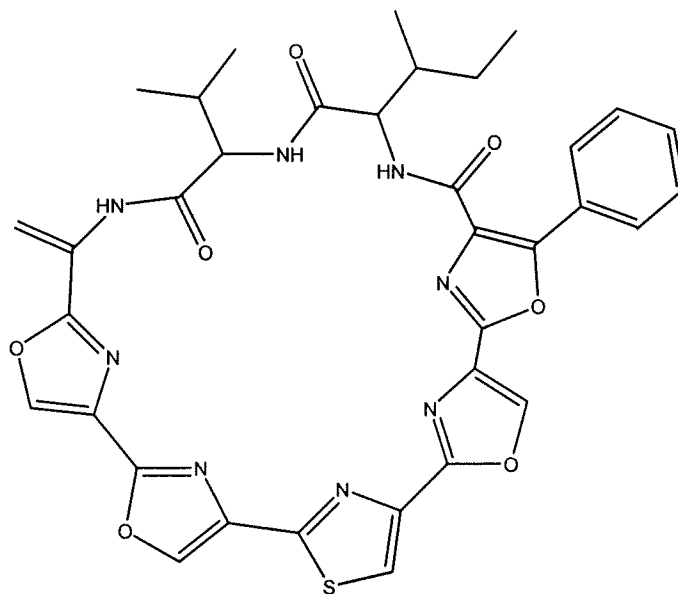
8. (Original) The compound according to claim 7, having the following stereochemistry



9. (Original) A process for producing a compound as defined in claim 1 which comprises synthesising a oxazole/thiazole/imidazole fragment, and introducing an aminoacidic fragment.

10. (Currently Amended) A process for preparing a compound as defined in claim 1 which comprises (i) cultivating a microorganism strain of the *Thermoactinomyces* genus a ~~microorganism~~ capable of producing it and (ii) isolating said compound from the cultured broth.

11. (Currently Amended) A process according to claim 10, wherein the compound isolated ~~prepared~~ is IB-01211 of formula:



16. **(Cancelled)**

17. (Currently Amended) A method of treatment of cancer which comprises administering an effective amount of a compound as defined in ~~any of claims~~ claim 1 to 8, or a pharmaceutically acceptable salt, derivative, prodrug or stereoisomer thereof.

18. (New) The process of claim 10, wherein the cultivating is performed in an aqueous nutrient medium with assimilable carbon and nitrogen sources and salts.